

HEPARIN SODIUM - heparin sodium injection, solution
Cantrell Drug Company

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Heparin Sodium 4,000 USP Units Added to 0.9% Sodium Chloride 1,000 mL Bag

HEPARIN

Sodium

4,000

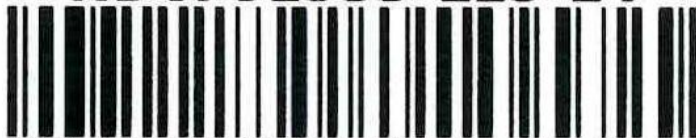
USP Units

Added to 0.9% Sodium Chloride 1,000 mL* Bag

(4 USP units/mL) *Volume & Concentration Exclude Manufacturer Overfill
Store at Room Temperature. Single-Dose Bag.
Hospital/Office Use Only. Injection Solution For IV Use.

**HIGH
ALERT**

NDC: 52533-125-24



(01) 0 0352533 12524 9



Rx Only

Each mL Contains: Heparin Sodium 4 USP units, Sodium Chloride 9.034 mg, trace amount of Benzyl Alcohol. pH adj: HCl/NaOH.
Outsourced Compounded Drug. Not for Resale.

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CANTRELL DRUG COMPANY

7321 Cantrell Road Little Rock, AR 72207
(877) 666-5222 www.cantrelldrug.com

LOT: xxxxxx

BUD:

CMPD Date: 03/13



HEPARIN SODIUM

heparin sodium injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-125
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	4 [USP U] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Sodium Chloride (UNII: 451W47IQ8X)	9 mg in 1 mL
BENZYL ALCOHOL (UNII: LKG8494WBH)	0.00004 mL in 1 mL
Water (UNII: 059QF0KO0R)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	HYDROCHLORIC ACID (UNII: QTT17582CB)	
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-125-24	1000 mL in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/09/2012	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014

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